25. (New) The method according to claim 1 wherein treatment of the cell population with the two or more immunotoxins causes low toxicity to CD34+ cells in the population.

#### REMARKS

The specification has been amended to correct typographical errors. No new matter has been introduced by these amendments. Claim 1 has been amended and new claims 15-25 have been added to more clearly define the invention. Upon entry of this amendment, Claims 1, 3, 6-8, 13, and 14-25 are pending in the application. Amendments to the claims have been made to place the claims in condition for allowance, or in a better form for appeal. Therefore, entry of this amendment is respectfully requested. A list of the pending claims are provided to the examiner for her convenience.

The previously amended Table 4, as detailed in the applicant's response to the Office Action mailed 7 August 2000 (Paper No 15), contains a clerical error. The amended Table 4 was intended to retain the data in the far right column indicating 1.0 µg/ml which shows a 2.5 log cell kill, not 0.1 µg/ml as indicated in the applicant's first response. It would have been obvious to one of ordinary skill in the art that, in light of the written specification, in particular Figure 1, that 0.1 µg/ml as detailed in Figure 4 was an error. Therefore, the amended Table 4 does not introduce new matter into the disclosure of the invention and the applicant respectfully requests removal of the objection under 35 U.S.C. § 132.

# Claim Rejections - 35 U.S.C. § 112

Claims 1, 3 and 6-8 remain rejected under 35 U.S.C. 112, first paragraph for reasons set forth in Paper No. 13, Section 6, pages 3-6. The Office Action (Paper No. 16) states, "The [Applicant's previous] arguments have been considered but have not been found persuasive (a') for the reasons previously set forth drawn to the unpredictability of anticancer drug discovery and differences between in vitro assays and complex conditions of in vivo therapy." Applicants respectfully traverse the rejection.

Applicants respectfully assert that the probability of successfully killing maliganant cells in vivo using the treatment as established in vitro is very high due to the 1) high specificity of the

antibodies, 2) the effectiveness of the combined immunotoxin approach 3) the low toxicity of the combined immunotoxin approach for non-maligant cells and, 4) previous data indicating the specificity of clinical localization of the Moc 31 antibody to tumor tissue compared to healthy tissue. The Applicant has demonstrated that the immunotoxin has a highly specific toxic effect on cancer cell lines but not on normal hematopoietic cells, including early progenitor cells. The early progenitor cells are highly sensitive to toxic effects of anticancer compounds, such as toxins, and the selective effect as demonstrated in the Applicant's current immunotoxin approach strongly argues for the same successful and selective effect in vivo.

It is generally accepted in the art that even though the transition from ex vivo to in vivo is complex, as indicated in the office action, this is the <u>only</u> accepted method for testing medicaments. Thus it is a general opinion in the art that it is not possible to test the effect of cancer medication, without a basis in ex vivo experiments as has been and is being done in large governmental and company drug screening programs.

Applicant respectfully requests reconsideration of this argument and request withdrawl of this rejection.

Claims 3 was rejected for the reasons previously set forth in Paper No. 13, Section 7, pages 6-9. The Office Action (Paper No. 16) indicates that neither antibody BM7 or MOC 31 are commercially available, are irrevocably available to the public with out restriction, or will be available if viable sample cannot be dispensed by S. Kaul or MCA Development, respectively.

Applicants provide information regarding the commercial availability of the BM7 and MOC 31 antibodies.

The monoclonal antibody BM7 can be obtained from:

MEDAC GmbH

Postfach 303629

D-20312 HAMBURG GERMANY

tel: 040/350920-0

fax: 040/350902-61

A description of the BM7 m-moclonal antibody can be found at www.medac.dc under "product portfolio" -> "diagnostics" -> "products" -> "oncology". A description sheet for the BM7 antibody is enclosed.

The MOC-31 antibody can be obtained from:

Zymed Laboratories, Inc.

458 Carlton Court

South San Francisco, CA 94080

tel.: 800.874.4494

fax: 650.871.4499

The MOC-31 catalog number is 18-0270; mouse anti-EGP-2 IgG<sub>1</sub>. A specification sheet for the MOC-31 antibody is enclosed.

Applicant respectfully requests reconsideration of this information and request withdrawl of this rejection.

#### Claim Rejections - 35 U.S.C. § 103

Claims 1 and 14 remain rejected under 35 USC 103 for the reasons previously set forth in Paper No. 13, Section 10, pages 12-15. The Office Action (Paper No. 16) indicates that the Applicant argues the references individually without clearly addressing the combined teachings. Applicants respectfully traverse the rejection.

The Applicant respectfully disagrees with the opinion of the Office Action and argues that none of the references alone or in combination teach or suggest the present invention, and one of skill in the art would not have looked to the cited references in attempting to derive the claimed invention.

The '254 patent teaches a DNA sequence encoding a GA 733.1 antigen recognized by a GA733 antibody. However, the GA 733.1 DNA sequence, following expression, can be post-translationally modified by glycosylation such that many epitopes are created and therefore the GA 733.1 antigen becomes nonspecific compared to the specific epitope recognized by the MOC 31 MAb. Therefore, the '254 patent is deficient and fails to teach to the current invention.

Second, it is not possible to use the combined teaching of the cited publications. The following essential features and requirements of the claimed invention are not met. First, the present method requires that the correct antibody is bound to the correct antigen and antigen epitope. This is essential both to obtain the required specificity and the activity of the immunotoxin. The combined publications fail to teach this.

Second, the antibodies <u>must</u> be connected to the optimal toxin that is antigen/antibody dependent. This is absolutely necessary in order to obtain the optimal cellular internalization and subsequent high activity. The combined publications also fail to teach this. For example, Lemoli's toxin would not be effective connected to the present antibody. The teaching of Bjorn together with Lemoli does not suggest the claimed invention due to the rigid requirements regarding the specificity of the claimed method.

Thirdly, all of the cited publications concern ex vivo methods and the combination of these fail to teach which antibodies and toxins can be connected to obtain the desired effect, or how these antibody/toxin combinations will work in vivo. In the claimed invention the immunotoxin is not affecting healthy tissue but is concentrated in tumor tissue. This is not suggested in the cited publications in combination.

It is known that the antibody is specific in that it will bind to antigens on cancer cells, but the combination of the cited publications did not suggest if the binding of the toxin to the antibody destroyed the antigen-antibody binding capacity. This had to be tested by experimentation.

Applicant respectfully requests consideration these arguments and requests withdrawl of this objection.

### New Grounds of Objection - Specification

The Office Action (Paper No. 16) indicates that the amendment filed August 14, 2000 is objected to under 35 U.S.C. 132 because it introduces new matter. Specifically, the Office Action states, "The amended Table 4 replaces the original 1.0 ug/ml with 0.1 ug/ml, retaining the 2.5 log cell kill. There is no supporting the specification as originally filed for a 0.1 ug/ml with a 2.5 log cell kill."

As addressed by the Applicant previously, regarding the Amendment in the specification of Table 4, the 0.1 ug/ml was a result of a clerical error and the amended Table 4 corrects this problem. Applicant has withdrawn Amended Table 4 as submitted in the papers filed 14 August, 2000. Applicant respectfully requests reconsideration of this amendment and requests withdrawl of this objection.

### New Grounds of Rejection - Claim Rejections - 35 U.S.C. § 112

The Office Action (Paper No. 16) indicates that Claims 1, 3, 6-8 and 14 are rejected under 35 U.S.C. 112 for lack of written description. Specifically, the Office Action states, "The limitation of active toxin fragments has no clear supporting the specification and the claims as originally filed." Furthermore, the Office action states, "...because claim 1 recites the phrase, "active toxin fragments". It is confusing because it is not clear what type of activity is being claimed."

The Applicant has removed "active" from Claim 1, as detailed in the Amendment.

Applicant respectfully requests reconsideration of this amendment and requests withdrawl of this rejection.

## CONCLUSION

Applicants respectfully assert that the claims 1, 3, 6-8, 13, and 14-25, upon entry of this amendment, are in a condition for allowance, and carnestly solicit a notice to that effect.

Applicants believe all of the outstanding objection and rejections have been addressed. If the Examiner has any questions regarding the foregoing, it is respectfully requested that she call the undersigned.

Respectfully Submitted,

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Date 1/30/2001

John J. Zresens Reg. No. 33,112 JJG/PLW/lc

Enclosures: Antibody specification sheets; Claims pending upon amendment entry.

KACLIENTS.07/07885/55USWO\d-Amendidoc

S/N 09/125,751

### Pending Claims Upon Entry of the Amendment Filed 03 January, 2001

- 1. (Thrice amended) Method to kill breast cancer cells or other carcinoma cells expressing target antigens in a cell population selected from the group consisting of cells comprising nucleated cells in peripheral blood and bone marrow cells comprising CD-34<sup>-1</sup> cells selected from the above nucleated cells, wherein the cell population is exposed to a combination of two immunotoxins, wherein each immunotoxin is composed of a conjugate between an antibody and a cell toxin, antigen binding antibody fragments and toxin fragments, or recombinantly produced antibodies, toxins, immuniotoxins or fragments thereof, wherein the antibodies are directed to epitopes on the antigen EGP2 expressed by the gene GA733-2 and to epitopes on the antigen expressed by the MUC1 gene and the toxin is Pseudomonas exotoxin A.
- 3. (Amended) Method according to claim 1, characterized in that the used antibodies are MOC31 and 595A6, or antigen binding fragments thereof.
- 6. (Amended) Method according to claim 1 wherein said exposure consists of administering the specific immunotoxins in vivo.
- 7. Method according to claim 6, characterized in that the immunotoxins are administered systemically, especially in case of malignant spread to tissues such as bone and bone marrow.
- 8. Method according to claim 6, characterized in that the immunotoxins are administered directly into the tumor or in the pleural and abdominal cavities.
- 14. (Amended) The method of claim 1, wherein said exposure consists of administering the immunotoxins ex vivo.

15.(New) A method for killing malignant cells in a cell population, the method comprising

obtaining the population of cells ex vivo that contains the malignant cells;

contacting the population of cells with at least two immunotoxins, wherein a first immunotoxin comprises a PE molecule conjugated to an antibody or an antibody fragment capable of binding an EGP2 antigen which is expressed by a GA733-2 gene and a second immunotoxin comprising a PE molecule conjugated to an antibody or an antibody fragment capable of binding an antigen encoded by the MUC1, MUC2, or MUC3 gene.

- 16. (New) The method according to claim 15, wherein the first immunotoxin comprises a PE molecule conjugated to a MOC31 antibody or an antigen-binding antibody fragment thereof, and the second immunotoxin comprises a PE molecule conjugated to a 595A6 antibody or an antigen-binding antibody fragment thereof.
- 17. (New) The method according to claim 15, wherein the cell population is obtained ex vivo from a cancer patient.
- 18. (New) The method according to claim 17, wherein the cell population comprises peripheral blood cells or bone marrow cells.
- 19. (New) The method according to claim 18, wherein the cell population comprises CD34+ cells
- 20. (New) The method according to claim 19, wherein the cell population is enriched or positively selected for CD34+ cells.
- 21. (New) The method according to claim 1 wherein treatment of the cell population with the two or more immunotoxins causes low toxicity to CD34+ cells in the population.

22. (New) A method for killing malignant cells in a patient, the method comprising

admistering to the patient a therapeutically effective amount of at least two immunotoxins, wherein a first immunotoxin comprises a PE molecule conjugated to an antibody or an antibody fragment capable of binding an EGP2 antigen which is expressed by a GA733-2 gene and a second immunotoxin comprises a PE molecule conjugated to an antibody or an antibody fragment capable of binding an antigen encoded by the MUC1, MUC2, or MUC3 genes.

- 23. (New) The method according to claim 22, wherein the patient is a cancer patient.
- 24. (New) The method according to claim 22, wherein the malignant cells are carcinomas.
- 25. (New) The method according to claim 1 wherein treatment of the cell population with the two or more immunotoxins causes low toxicity to CD34+ cells in the population.